<sup>100</sup> 
$$MHy^{+} + H_20 \rightleftharpoons NH_3 + H_3 o^{+}$$
;  $K_a = [NH_3] [H_3o^{+}]_{=} = 5,56 \times 10^{-10}$   
Now that we have Ka, solve as a normal weak acid.  
Species [Jnitial]  $\Delta$  [Equilibrium]  
 $H_30^{+}$   $O$   $+X$   $X$   
 $NH_3$   $O$   $+X$   $X$   
 $NH_4^{+}$   $O,100 - X$   $O,100 - X$   
 $(x)(x) = 5.56 \times 10^{-10}$   $X = 7.45 \times 10^{-6} = [H_3o^{+}]$   
 $\downarrow \chi^{22} = 5.56 \times 10^{-10}$   $X = 7.45 \times 10^{-6} = 5.12$   
Compare:  
 $PH = -log_{10}(7.45 \times 10^{-6}) = 5.12$   
Compare:  
 $PH of 0.100 M nitrous acid: 2.17$   
 $PH of DI water: 7.00$ 

O,100M NaC2H302, Find PH  $Na(2H_2O_2 \rightarrow Na^+ + C_2H_3O_2^-)$  $\mathcal{N}_{A}^{+}$  Not a proton donor (acid), since no H+ to donate. Also not likely to be basic, as the positive charge should repel H+. NEUTRAL.  $(2H_3O_2^-:$  Negative charge ... this might accept protons and be a base. If so, the reaction looks like this:  $C_2H_3O_7 + H_2O \rightleftharpoons H(2H_3O_2 + OH)$  $\angle$  Stable in water? This is ACETIC ACID, a common WEAK ACID. Weak acids are stable in water, since very few weak acid molecules ionize! So, acetate ion is a base. (2H30, + H20 = H(2H302 + 0H - ; Kb = [H(2H302][0H-]] ->  $[(_{1}H_{1}O_{1}^{-}]]$ Appendix I doesn't list a Kb for acetate ion, but Appendix H has a Ka for acetic acid, its conjugate. Ka, H(2Hjuz = 1.8×10-5 KaxKb = 1.0×10-14 (1.8×10-5) Kb = 1.0×10-14  $K_{\rm b} = 5.56 \times 10^{-10}$ 

<sup>135</sup>  

$$C_{2}H_{3}O_{2}^{-} + H_{2}O \rightleftharpoons H(2H_{3}O_{2} + OH^{-1}) K_{0} = \frac{[H(_{2}H_{3}O_{2}][OH^{-1}]}{[C(_{2}H_{3}O_{2}^{-1}]} = 5.56 \times 10^{-16}$$
Once we find Kb, just solve like any other weak base.  

$$\frac{Specres}{OH^{-1}} \frac{[E_{1}n_{1}f_{1}M_{1}]}{OH^{-1}} \bigtriangleup \frac{[E_{1}n_{1}f_{1}h_{1}]}{[C(_{2}H_{3}O_{2}^{-1}]} \qquad \text{Let "x" equal the change in hydroxide ion concentration!}$$

$$\frac{H(_{2}H_{3}O_{2})}{(2H_{3}O_{2})} \frac{O_{1}100}{O_{1}100} + X \times \frac{X}{(2H_{3}O_{2})} O_{1}100} + X \otimes \frac{X}{(X)(X)} = 5.56 \times 10^{-16}$$

$$\int_{(0,100-X)}^{O(100-X)} O_{1}100 - X = 0.100 + X \otimes \frac{100}{(X \times 10^{-5})} = 4.12$$

$$PUH = -10g_{10}(2.45 \times 10^{-5}) = 4.12$$

$$PH = POH = 140,00$$

$$PH = 140,00$$

For comparison:

0.100 M sodium acetate, pH = 9.87

0.100 M ammonia, pH = 11.13

0.100 M NaOH (strong base), pH = 13.00

The acetate ion is basic, but it's a very weak base!

$$O.100 M NaCl, Find pH$$
  
 $NaCl \rightarrow Na^+ + Cl^-$ 

140

- $\mathcal{N}_{\alpha}^{+}$ : Not a proton donor (acid), since no H+ to donate. Also not likely to be basic, as the positive charge should repel H+. NEUTRAL.
- C Not a proton donor (acid), since it has no protons (H+) to donate. It does have a negative charge, and may attract protons. If so, this would be the reaction:

$$(1^{-} +H_2 O \rightleftharpoons HCI + OH^{-})$$

This is hydrochloric acid, a STRONG ACID. Strong acids completely ionize, so they are NOT stable in water. This reaction won't happen! Chloride ion should be NEUTRAL.

In a NaCl solution, water itself controls the pH. So, the pH will be 7.00

## <sup>141</sup> POLYPROTIC ACIDS

... what's special about phosphoric acid?

 $K_{a1} = 7.5 \times 10^{-3}$   $K_{a2} = 6.2 \times 10^{-8}$  $K_{a3} = 4.2 \times 10^{-13}$ 

(1) 
$$H_3 PO_4 + H_2 O \rightleftharpoons H_2 PO_4^- + H_3 O^+$$
  
(2)  $H_2 PO_4^- + H_2 O \rightleftharpoons H PO_4^{2-} + H_3 O^+$   
(3)  $H PO_4^{2-} + H_2 O \rightleftharpoons PO_4^{3-} + H_3 O^+$ 

Phosphoric acid has THREE acidic protons!

The first dissocation is dominant here, and for simple calculations of phosphoric acid in water, we will simply use the first ionization and ignore the other two.

Remember: This is a weak acid. It exists in water mostly as undissociated phosphoric acid molecules.

Solve the equilibrium of phosphoric acid's FIRST proton:  
H<sub>3</sub> PO<sub>44</sub> + H<sub>2</sub>O 
$$\rightleftharpoons$$
 H<sub>2</sub> PO<sub>47</sub> + H<sub>3</sub>O<sup>+</sup>; K<sub>6</sub> = 7.5 × 10<sup>-3</sup>  
K<sub>6</sub> =  $\frac{1}{2}$  H<sub>2</sub> PO<sub>47</sub> = 7.5 × 10<sup>-3</sup>  
K<sub>6</sub> =  $\frac{1}{2}$  H<sub>3</sub> PO<sub>4</sub> = 7.5 × 10<sup>-3</sup>  
Let "x' equal the change in hydronium ion concentration  
H<sub>3</sub>O<sup>+</sup> O + X ×  
H<sub>2</sub> PO<sub>4</sub> O + X ×  
H<sub>2</sub> PO<sub>4</sub> O + X ×  
(X)(X) = 7.5 × 10<sup>-3</sup>  
(V, 1(X) = 7.5 ×

<sup>143</sup> Find the pH of a solution prepared by dissolving 3.00 g of ammonium nitrate (FW=80.052 g/mol) solid into enough water to make 250. mL of solution.

$$\begin{aligned} NH_{\mathcal{Y}} NO_{3} & \longrightarrow NH_{\mathcal{Y}}^{+} + NO_{3}^{-} \text{ Is this salt acidic, basic, or neutral?} \\ NH_{\mathcal{Y}}^{+} \text{; Has protons? Might be a proton donor (acid):} \\ NH_{\mathcal{Y}}^{+} + H_{2}O & \xrightarrow{NH_{3}} + H_{3}O^{+} \\ \hline L \text{ Ammonia. A WEAK BASE! This is stable in water, so} \\ \text{this reaction will go. Ammonium ion is ACIDIC.} \\ NO_{3}^{-} \text{; Could this be a proton acceptor (base)?} \\ NO_{3}^{-} + H_{2}O & \xrightarrow{H_{N}O_{3}} + 0H^{-} \\ \hline L \text{ Nitric acid, a STRONG ACID. Not stable in water, so this} \\ \text{reaction doesn't go! Nitrate is NEUTRAL.} \\ NH_{\mathcal{Y}}^{+} + H_{2}O & \xrightarrow{H_{3}} + H_{3}O^{+} \text{; } K_{4} = \frac{(NH_{3})[H_{3}O^{+}]}{[NH_{4}^{+}]^{2}} = \frac{7}{2} \\ \text{Appendix I has Kb for ammonia, so we can calculate Ka for ammonium ion:} \\ Kb_{1}NH_{3} = 1.8 \times 10^{-5} \text{ Ka x Kb} = 1.0 \times 10^{-14} \\ \text{Ka} (1.8 \times 10^{-5}) = 1.0 \times 10^{-14} \text{; } K_{4} = 5.5 \text{ b x lo}^{-15} \\ \text{We'll need to know the MOLAR concentration of the solution. Calculate.} \\ 3 \text{ $DO_{6}NH_{4}NO_{3} \times \frac{MO_{1}NH_{4}NO_{3}}{(O.052 \text{ g}NH_{4}NO_{3})}} = 0.1049025(33 \text{ M} NH_{4}NO_{3}) \\ O.250 L \\ \Lambda = \frac{0.037475(406 \text{ mu})}{(250 \text{ c})} = 1.0 \times 10^{-14} \text{ M}_{4}NO_{3} = 0.1049025(33 \text{ M} NH_{4}NO_{3}) \\ \end{array}$$

$$NHy^{+} + H_2 O = NH_3 + H_3 O + K_4 = \frac{[NH_3][H_3 O^{+}]}{[NH_4^{+}]} = 5.56 \times 10^{-10}$$

in

0.1499025633 M NH4 NO3

Solve the acid ionization equilibrium.  
Species [Initial] 
$$\Delta$$
 [Cequilibrium]  
H<sub>30</sub>+ 0 + x x  
NH<sub>3</sub> 0 + x x  
NH<sub>3</sub> 0 + x x  
NH<sub>4</sub>+ 0.149902563 - X 0.1499025637-X  
(x)(x) = 5.56 x10<sup>-10</sup>  
(0.1499025637-X) = 5.66 x10<sup>-10</sup>  
xLL0.150,50  
-x - 0,150  
x<sup>2</sup>  
0.1499025637 = 5,56 x10<sup>-10</sup>  
x = 9.12939 x10<sup>-6</sup> =  $\Sigma H_30^{+3}$ 

144